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<b>(21) International Application Number:</b> PCT/US95/04045 <b>(22) International Filing Date:</b> 13 April 1995 (13.04.95) <b>(30) Priority Data:</b> 08/227,961 ) 15 April 1994 (15.04.94) US <b>(71) Applicant:</b> METASYN, INC. [US/US]; 71 Rogers Street, Cambridge, MA 02142-1118 (US). <b>(72) Inventors:</b> SCOTT, Daniel, M.; 42 Nylander Way, Acton, MA 01720 (US). LAUFFER, Randall, B.; 23 Sumner Road #2, Brookline, MA 02146 (US). <b>(74) Agents:</b> HALEY, James, F., Jr. et al.; Fish & Neave, 1251 Avenue of the Americas, New York, NY 10020-1104 (US).		<b>(81) Designated States:</b> AM, AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LT, LU, LV, MD, MG, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TT, UA, UG, UZ, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG), ARIPO patent (KE, MW, SD, SZ, UG).  <b>Published</b> <i>With international search report.</i>
<b>(54) Title:</b> CONTRAST AGENTS FOR DIAGNOSTIC IMAGING  <b>(57) Abstract</b>  Paramagnetic metal complexes of 1-(p-n-butylbenzyl)diethylenetriamine pentaacetic acid and pharmaceutically acceptable salts thereof are effective as tissue-specific MRI contrast agents, and in particular as hepatobiliary MRI contrast agents.		

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CONTRAST AGENTS FOR DIAGNOSTIC IMAGINGTECHNICAL FIELD OF THE INVENTION

The present invention relates to diagnostic  
5 nuclear magnetic resonance imaging (MRI or MR imaging).  
In particular, this invention relates to novel  
compounds that upon administration to a patient, can  
enhance the contrast in images of the patient's organs  
and tissue obtained through MRI. This invention also  
10 relates to pharmaceutical compositions comprising these  
compounds and to methods of making and using the  
compounds and compositions for MRI contrast  
enhancement.

BACKGROUND OF THE INVENTION

15 Nuclear magnetic resonance imaging (MRI) is  
one of the most useful diagnostic tools in medicine  
today. The technique is based on the discovery that  
atomic nuclei possess a magnetic moment which can be  
detected using a magnetic field. When placed in a  
20 magnetic field, nuclei arrange themselves in a manner  
that aligns their magnetic moments either with or  
against the direction of the magnetic field. Those  
that align against the field (so-called  $\beta$ -spin nuclei)  
are higher in energy than those aligned with the field  
25 (so-called  $\alpha$ -spin nuclei). The energy difference  
between  $\alpha$ - and  $\beta$ -spin nuclei is directly proportional  
to the strength of the applied field. To measure this

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difference in energy, the nuclei are subjected to electromagnetic radiation. Low energy  $\alpha$ -spin nuclei absorb electromagnetic radiation causing them to "flip" against the field (or out of alignment with the field) into a high energy  $\beta$ -spin. The frequency of the absorbed radiation, known as the "resonance frequency," reveals the  $\alpha$ - $\beta$  energy difference. Alternatively, high energy  $\beta$ -spin nuclei or non-aligned nuclei may "relax" into a low energy  $\alpha$ -spin state with concomitant release of electromagnetic radiation. Both the resonance frequency and the relaxation rates (measured in terms of relaxation rate  $(1/T_1)$  wherein  $T_1$  = the time that it takes for non-aligned nuclei to relax) for any nucleus is dependent on the magnetic environment surrounding that nucleus. For example, nuclei within a substantial (strong) electron shell are shielded from external magnetic fields and as a result possess a smaller  $\alpha$ - $\beta$  energy difference, compared to nuclei within a weak electron shell. Also, nuclei situated in the proximity of other paramagnetic nuclei relax faster and have higher relaxation rates than those for which such a relaxation mechanism is not available.

It is the difference in the nuclear magnetic relaxation rates of water protons that is most commonly used as the source of information in diagnostic MRI. Water is ubiquitously found in soft tissue. The hydrogen atoms in water possess paramagnetic nuclei (protons) that, when subjected to resonance frequency and relaxation-rate measurements, provide direct information regarding their physical microenvironment. The information can then be processed through computational techniques to obtain detailed anatomical images. Inspection of these images can reveal the presence of diseased or abnormal tissue, lesions and fractures or other malfunctions that may be present.

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Not surprisingly, MR imaging of some tissue is more difficult than others. This may be due to any number of factors which result in low contrast and hence lack of resolution in the MR images obtained.

- 5 Previous efforts to overcome such problems have led to the discovery of so-called contrast enhancement agents. Typically, such agents contain a paramagnetic metal ion which is capable of altering the relaxation rates of water protons in its proximity. Upon administration to
- 10 a patient, the metal-containing contrast agent is absorbed by various organs depending on the patient's metabolic and excretion pathways. Once absorbed, the contrast agent alters the relaxation rates of water protons in the organ or tissue in which it resides.
- 15 The MR images of that organ or tissue thus achieve enhanced contrast with respect to neighboring tissue which contains lower concentrations or none of the paramagnetic agent.

- Because of the acute toxicity of most
- 20 paramagnetic metals, however, ordinary inorganic salts of paramagnetic metals are unsatisfactory as contrast agents. A solution to this problem is to use an organic chelating ligand or metal-sequestering agent. Through complexation to the metal, the organic
- 25 chelating ligand would prevent release of free, toxic metal yet allow proton relaxation enhancement by acting as a non-toxic paramagnetic carrier.

- To be effective, MRI contrast agents (paramagnetic metal-ligand complexes) must satisfy
- 30 several criteria. They must be stable and have high formation constants so that release of the toxic metal is prevented. They must be sufficiently soluble in aqueous solutions to facilitate their administration to a patient. And, they must be capable of efficient
- 35 enhancement of relaxation rates of water protons in

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solution. Efficiency is generally measured in terms of "relaxivity", which is defined as the increase in relaxation rate per concentration of the paramagnetic complex measured in units of mM.

5           Gries et al. have described complexes for use as diagnostic agents in United States patent 4,647,447. Also, the active paramagnetic ingredient of the FDA-approved MRI contrast agent Magnevist® is a complex of diethylenetriaminepentacetic acid and gadolinium (III).  
10 In United States patent 4,899,755, Lauffer and Brady describe how paramagnetic metal-ligand complexes can be designed and synthesized to target specific tissue for MRI enhancement. This tissue-specific approach provides several improvements over the previously  
15 reported non-specific methods. Qualitatively, tissue-specific agents provide better MR images of the targeted tissues such as the liver and the bile duct. Quantitatively, tissue-specific contrast agents can be used in lower concentrations to achieve image  
20 enhancements similar to that observed with non-specific agents at higher doses. Thus, previously undetectable (or difficult to detect) liver tumors or malfunctions in the biliary system can be detected using hepato-biliary-specific contrast agents.

25           Another improvement upon the non-specific methods of the art is provided in United States patent 4,880,008 where the relaxivity-enhancement of contrast agents is shown to improve upon noncovalent binding to specific tissue proteins. This strategy not only  
30 allows targeting of specific tissue containing the binding proteins, it also provides image enhancement at even lower doses than previously possible (due to improved relaxivity enhancement of the protein-bound agents).

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In spite of these advances, there remains a need for MRI contrast agents that combine favorable stability, solubility, relaxivity-enhancement and protein-binding characteristics with a tissue-specific pharmacokinetic profile.

#### SUMMARY OF THE INVENTION

The present invention provides improved compounds that are useful as tissue-specific MRI contrast agents, compositions containing those compounds and methods for using them. In particular, the present invention provides paramagnetic metal complexes of 1-(p-n-butylbenzyl)diethylenetriamine pentaacetic acid and pharmaceutically acceptable salts thereof as useful and efficient agents for improved tissue-specific enhancement of contrast in MRI of mammalian organs and tissues.

#### DETAILED DESCRIPTION OF THE INVENTION

The tissue-specific contrast agents of this invention comprise complexes of a paramagnetic metal and 1-(p-n-butylbenzyl)diethylenetriamine pentaacetic acid and pharmaceutically acceptable salts thereof. The stereochemistry at position 1 may be R or S, and the paramagnetic complex may be used in racemic or enantiomerically pure form. Preferred paramagnetic metals of this invention are selected from the group consisting of Gd(III), Fe(III), Mn(II and III), Cr(III), Cu(II), Dy(III), Tb(III), Ho(III), Er(III) and Eu(III). The most preferred paramagnetic metal is Gd(III).

Pharmaceutically acceptable salts of this invention include those derived from inorganic or organic acids and bases. Included among such acid salts are the following: acetate, adipate, alginate,

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aspartate, benzoate, benzenesulfonate, bisulfate, butyrate, citrate, camphorate, camphorsulfonate, cyclopentanepropionate, digluconate, dodecylsulfate, ethanesulfonate, fumarate, glucoheptanoate, glycerophosphate, hemisulfate, heptanoate, hexanoate, hydrochloride, hydrobromide, hydroiodide, 2-hydroxyethanesulfonate, lactate, maleate, methanesulfonate, 2-naphthalenesulfonate, nicotinate, oxalate, pamoate, pectinate, persulfate, 3-phenylpropionate, picrate, pivalate, propionate, succinate, tartrate, thiocyanate, tosylate and undecanoate. Base salts include ammonium salts, alkali metal salts, such as sodium and potassium salts, alkaline earth metal salts, such as calcium, magnesium and zinc salts, salts with organic bases, such as dicyclohexylamine salts, N-methyl-D-glucamine, and salts with amino acids such as arginine, lysine, and so forth. Also, the basic nitrogen-containing groups can be quaternized with such agents as lower alkyl halides, such as methyl, ethyl, propyl, and butyl chloride, bromides and iodides; dialkyl sulfates, such as dimethyl, diethyl, dibutyl and diamyl sulfates, long chain halides such as decyl, lauryl, myristyl and stearyl chlorides, bromides and iodides, aralkyl halides, such as benzyl and phenethyl bromides and others. Water or oil-soluble or dispersible products are thereby obtained. The preferred salts of this invention are the N-methyl-D-glucamine, calcium and sodium salts.

The pharmaceutical compositions of this invention comprise any of the complexes of the present invention, or pharmaceutically acceptable salts thereof, together with any pharmaceutically acceptable carrier, adjuvant or vehicle. Pharmaceutically acceptable carriers, adjuvants and vehicles that may be used in the pharmaceutical compositions of this



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invention include, but are not limited to, ion exchangers, alumina, aluminum stearate, lecithin, serum proteins, such as human serum albumin, buffer substances such as phosphates, glycine, sorbic acid, potassium sorbate, TRIS (tris(hydroxymethyl)amino-  
methane), partial glyceride mixtures of saturated vegetable fatty acids, water, salts or electrolytes, such as protamine sulfate, disodium hydrogen phosphate, potassium hydrogen phosphate, sodium chloride, zinc  
salts, colloidal silica, magnesium trisilicate, polyvinyl pyrrolidone, cellulose-based substances, polyethylene glycol, sodium carboxymethylcellulose, polyacrylates, waxes, polyethylene-polyoxypropylene-block polymers, polyethylene glycol and wool fat.

According to this invention, the pharmaceutical compositions may be in the form of a sterile injectable preparation, for example a sterile injectable aqueous or oleaginous suspension. This suspension may be formulated according to techniques known in the art using suitable dispersing or wetting agents and suspending agents. The sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example as a solution in 1,3-butanediol. Among the acceptable vehicles and solvents that may be employed are water, Ringer's solution and isotonic sodium chloride solution. In addition, sterile, fixed oils are conventionally employed as a solvent or suspending medium. For this purpose, any bland fixed oil may be employed including synthetic mono- or di-glycerides. Fatty acids, such as oleic acid and its glyceride derivatives are useful in the preparation of injectables, as are natural pharmaceutically-acceptable oils, such as olive oil or castor oil, especially in their polyoxyethylated

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versions. These oil solutions or suspensions may also contain a long-chain alcohol diluent or dispersant, such as Ph. Hely or similar alcohol.

The compounds and pharmaceutical compositions  
5 of the present invention may be administered orally, parenterally, by inhalation spray, topically, rectally, nasally, buccally, vaginally or via an implanted reservoir in dosage formulations containing conventional non-toxic pharmaceutically-acceptable  
10 carriers, adjuvants and vehicles. The term "parenteral" as used herein includes subcutaneous, intravenous, intramuscular, intra-articular, intra-synovial, intrasternal, intrathecal, intrahepatic, intralesional and intracranial injection or infusion  
15 techniques.

When administered orally, the pharmaceutical compositions of this invention may be administered in any orally acceptable dosage form including, but not limited to, capsules, tablets, aqueous suspensions or  
20 solutions. In the case of tablets for oral use, carriers which are commonly used include lactose and corn starch. Lubricating agents, such as magnesium stearate, are also typically added. For oral administration in a capsule form, useful diluents  
25 include lactose and dried corn starch. When aqueous suspensions are required for oral use, the active ingredient is combined with emulsifying and suspending agents. If desired, certain sweetening, flavoring or coloring agents may also be added.

30 Alternatively, when administered in the form of suppositories for rectal administration, the pharmaceutical compositions of this invention may be prepared by mixing the agent with a suitable non-irritating excipient which is solid at room temperature  
35 but liquid at rectal temperature and therefore will

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melt in the rectum to release the drug. Such materials include cocoa butter, beeswax and polyethylene glycols.

As noted before, the pharmaceutical compositions of this invention may also be administered topically, especially when the target of treatment includes areas or organs readily accessible by topical application, including the eye, the skin, or the lower intestinal tract. Suitable topical formulations are readily prepared for each of these areas or organs.

10 Topical application for the lower intestinal tract can be effected in a rectal suppository formulation (see above) or in a suitable enema formulation. Topically-transdermal patches may also be used.

15 For topical applications, the pharmaceutical compositions may be formulated in a suitable ointment containing the active component suspended or dissolved in one or more carriers. Carriers for topical administration of the compounds of this invention include, but are not limited to, mineral oil, liquid petrolatum, white petrolatum, propylene glycol, polyoxyethylene, polyoxypropylene compound, emulsifying wax and water. Alternatively, the pharmaceutical compositions can be formulated in a suitable lotion or cream containing the active components suspended or dissolved in one or more pharmaceutically acceptable carriers. Suitable carriers include, but are not limited to, mineral oil, sorbitan monostearate, polysorbate 60, cetyl esters wax, cetearyl alcohol, 20 2-octyldodecanol, benzyl alcohol and water.

For ophthalmic use, the pharmaceutical compositions may be formulated as micronized suspensions in isotonic, pH adjusted sterile saline, or, preferably, as solutions in isotonic, pH adjusted sterile saline, either with or without a preservative

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such as benzylalkonium chloride. Alternatively, for ophthalmic uses, the pharmaceutical compositions may be formulated in an ointment such as petrolatum.

For administration by nasal aerosol or  
5 inhalation, the pharmaceutical compositions of this invention are prepared according to techniques well-known in the art of pharmaceutical formulation and may be prepared as solutions in saline, employing benzyl alcohol or other suitable preservatives, absorption  
10 promoters to enhance bioavailability, fluorocarbons, and/or other conventional solubilizing or dispersing agents.

The amount of active ingredient that may be combined with the carrier materials to produce a single  
15 dosage form will vary depending upon the host treated, the sensitivity of the MR imaging instrument, the target tissue of the imaging experiment, the particular mode of administration and the intended effect of use. It should be understood, however, that a specific  
20 dosage regimen for any particular patient will depend upon a variety of factors, including the activity (induced relaxivity) of the specific agent employed, the age, body weight, general health, sex, diet, time of administration, rate of excretion, drug combination,  
25 and the judgment of the treating physician.

Dosage levels of between about 0.1 and about 1000  $\mu\text{mol/kg}$  body weight per day, preferably between about 0.5 and about 300  $\mu\text{mol/kg}$  body weight per day of the active ingredient compound are useful. A typical  
30 preparation will contain between about 1-1000 mM of the active complex. Preferably, such preparations contain between about 10-500 mM of the active complex.

To preserve shelf stability, to enhance the safety of the contrast agents and to prevent in vivo  
35 release of toxic metal (or ligand), according to one

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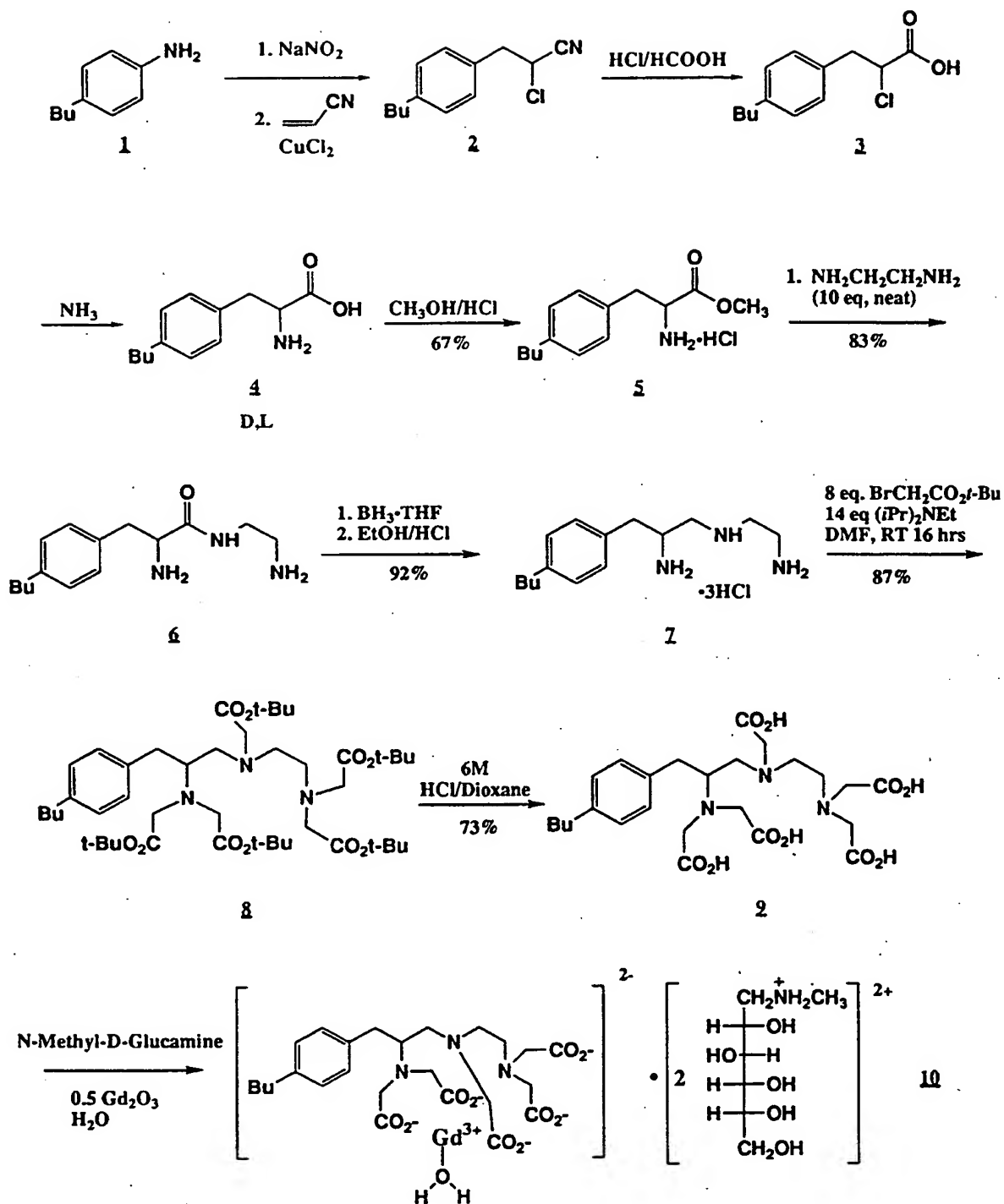
embodiment of this invention the compositions further contain a free organic ligand or an excess of 1-(p-n-butylbenzyl)diethylenetriamine pentaacetic acid and pharmaceutically acceptable salts of such ligands, for example, calcium, sodium and meglumine (N-methyl-D-glucamine) salts and combination salts such as a calcium sodium salt. The pharmaceutically acceptable compositions contain between 0-5% (molar ratio) of excess organic ligand or a pharmaceutically acceptable salt thereof, preferably 0-1%, and most preferably 0-0.5%. The use of compositions with an excess of a chelating ligand in diagnostic imaging has been disclosed in greater detail in International patent publication WO 90/03804, European patent publications EP-A-0463644 and EP-A-454078 and United States patent 5,078,986.

The tissue-specific contrast agents of this invention may be readily obtained using conventional techniques. Scheme 1 illustrates a representative example of a process for the preparation of gadolinium(III)-1-(p-n-butylbenzyl)diethylenetriamine pentaacetate dimeglumine salt (MS-264). It will be appreciated by those of skill in the art that the below synthetic processes are not intended as a comprehensive list of all means by which the contrast agents or the intermediates of this invention may be synthesized. Further methods or modifications of the below process will be evident to those of skill in the art.

Starting with 4-n-butylaniline, the process involves diazotization, followed by metal-catalyzed coupling with acrylonitrile or an equivalent (e.g., an alkyl ester of acrylic acid). The product is converted to p-n-butylphenylalanine which is then processed to obtain its ethylenediamine amide. Reduction of the amide to the amine using a known reducing agent,

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## Scheme 1



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preferably borane-THF complex, provides the p-n-butylbenzyl derivative of diethylenetriamine. The nitrogen atoms in this compound are alkylated using an alkyl ester of bromoacetic acid such as t-butyl-bromoacetate to give, after ester hydrolysis, the desired organic ligand.

Complexation to the paramagnetic metal can be achieved using any of the well-known procedures in the art. Scheme 1 provides an example of such a procedure involving the reaction of the ligand with gadolinium oxide in the presence of N-methyl-D-glucamine. Similar procedures using other paramagnetic metal oxides or salts instead of  $Gd_2O_3$  and other bases instead of N-methylglucamine provide further paramagnetic contrast agents of this invention.

Optically active contrast agents may also be prepared using, for example, optically active p-n-butylphenylalanine or using other optically active intermediates in the process illustrated in Scheme 1. Optically active intermediates can, in turn, come from resolution of a corresponding racemic mixture or from stereoselective synthesis. Methods for both resolution of racemic mixtures and stereoselective synthesis of optically active compounds are well known and require no further description here.

The contrast agents of this invention are surprisingly efficient and tissue-specific in MRI of mammalian tissue. Thus, this invention provides a method for tissue-specific contrast enhancement of MR images of mammalian organs and tissues. In particular, the method allows for contrast enhancement of MR images of the hepatobiliary system. For example, MS-264, the preferred contrast agent of this invention, exhibits surprisingly superior liver and bile specificity relative to structurally similar contrast agents

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previously disclosed (see Examples section). MS-264 can also be used for tissue-specific imaging of tumors and the blood pool.

Following administration of the contrast agent, NMR imaging is carried out; the choice of pulse sequence (inversion recovery, IR; spin echo, SE; gradient echo, GE; turbo FLASH,; etc.) and the values of the imaging parameters (echo time, TE; inversion time, TI; repetition time, TR; flip angle, etc.) will be governed by the diagnostic information sought. In general, if one desires to measure  $T_1$ , then TE should be less than 30 milliseconds (or the minimum value) to maximize  $T_1$ -weighting. Conversely, if one desires to measure  $T_2$ , then TE should be greater than 30 milliseconds to minimize competing  $T_1$  effects. TI and TR will remain approximately the same for both  $T_1$ - and  $T_2$ -weighted images; TI and TR are generally on the order of about 200-600 and 100-1000 milliseconds, respectively.

It is well known that ligands form complexes that are useful as MRI contrast agents may also be used in chelate complexes for diagnostic X-ray imaging. Therefore, according to another embodiment of this invention, metal complexes of 1-(p-n-butylbenzyl) diethylenetriamine pentaacetic acid and pharmaceutically acceptable salts thereof are used for diagnostic X-ray imaging.

In order that this invention may be more fully understood, the following examples are set forth. These examples are for the purpose of illustration only and are not to be construed as limiting the scope of the invention in any way.



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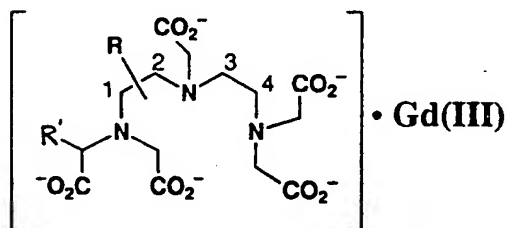
EXAMPLESRelaxivity in rat liver cytosol

Rat liver cytosol was prepared by homogenizing rat livers (25% w/v) in 0.25 M sucrose/0.01 M phosphate buffer, pH 7.4, centrifuging at 100,000 g for 90 min at 4°C, and concentrating the supernatant by pressure ultrafiltration to get the cytosol protein concentration close to that in vivo (5-10%). The relaxivity of the Gd(III) complexes in cytosol was measured at 37 °C using a Bruker 20 MHz Minispec equipped with a variable temperature probe. The inversion-recovery pulse sequence with 10 data points was used for  $T_1$  measurements.  $1/T_1$  relaxation rates of solutions with varying concentrations of the chelates (0.01-0.5 mM) were determined. The limiting slope of the  $1/T_1$  vs concentration plot at low concentrations of the chelates in units of  $s^{-1} \text{ mM}^{-1}$  was taken as the relaxivity (R1) of the chelate in cytosol. Results are summarized in TABLE 1 (Bn = benzyl).

Higher relaxivity values imply higher binding affinity to cytosolic proteins and/or greater relaxivity when bound.

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TABLE 1. Relaxivity In Rat Liver Cytosol



<u>DTPA derivative</u>		<u>R1 in liver cytosol</u> <u>20, MHz, 37°C</u> <u>(s<sup>-1</sup> mM<sup>-1</sup>)</u>
5	DTPA (R=R'=H)	7.1
	BOPTA (R=H, R'=BnOCH <sub>2</sub> )	11.3
<u>R (R'=H)</u>		
10	1-(benzyl)	11.7
	2-(benzyl)	nd
	1-(phenylethyl) [CH <sub>2</sub> CH <sub>2</sub> Ph]	13.6
	1-(p-ethoxybenzyl) [EOB-DTPA]	14.4
	1-(p-ethylbenzyl)	16.4
	1-(p-isopropylbenzyl)	18.7
	1-(p-n-butylbenzyl) [MS-264]	23.8
	1-(2,4-di-Me-benzyl)	15.3
	1-(2,4,6-tri-Me-benzyl)	17.9
	1,3-dibenzyl	23.7
20	1,4-dibenzyl	20.0
	1-(p-ethylbenzyl)-3-benzyl	nd
	1,3-di(p-ethylbenzyl)	nd
	1-(p-n-butylbenzyl)-3-benzyl	nd

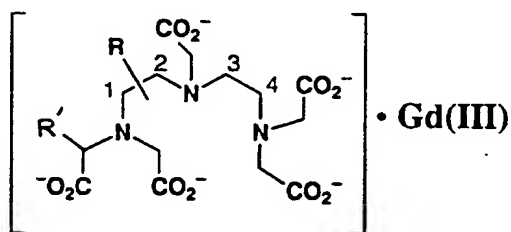
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Rat liver uptake and  $1/T_1$  relaxation rates of rat liver

Un-anesthetized adult male Sprague Dawley rats (200-300 g) were injected in the tail vein with 50  $\mu\text{mol/kg}$  of gadolinium chelate labeled with  $^{153}\text{Gd}$ .  
5 Rats (approx. 3 per time period) were euthanized by cervical dislocation (while anesthetized by  $\text{CO}_2$ ) at 1, 5 and 30 min. post-injection. The liver was analyzed for  $^{153}\text{Gd}$  concentration using a gamma counter. The percentage of the injected dose (% ID) in the liver was  
10 determined for each time point and the integrated area under the mean liver % ID time curve (AUC) was calculated.

For the 5 min. time period, the  $1/T_1$  proton relaxation rate of the excised liver was measured at  
15 20 MHz and 37 °C as described above. Higher  $1/T_1$  values are associated with greater MRI signal intensity on  $T_1$ -weighted images and are due to greater liver uptake of the agent and/or greater relaxivity in the micro-environments of the liver. Results are summarized in  
20 TABLE 2.

TABLE 2. Uptake and Induced Proton Relaxation Rates in Rat Liver



		Area under the % inj. dose in rat liver vs time curve (AUC) from 0 to 30 min*	1/T1 of excised rat liver 5 min. post-inj. (s <sup>-1</sup> ; 20 MHz, 37°C)
5	<u>DTPA derivative</u>		
	DTPA (R=R'=H)	35 (n=9)	4.64 ± 0.16 (n=3)
	BOPTA (R=H, R'=BnOCH <sub>2</sub> )	nd	
	<u>R (R'=H)</u>		
10	1-(benzyl)	213 (n=9)	nd
	2-(benzyl)	nd	nd
	1-(phenylethyl) [CH <sub>2</sub> CH <sub>2</sub> Ph]	nd	nd
	1-(p-ethoxybenzyl) [EOB-DTPA]	408 (n=9)	10.1 ± 1.5 (n=3)
	1-(p-ethylbenzyl)	469 (n=10)	12.3 ± 0.9 (n=3)
	1-(p-isopropylbenzyl)	400 (n=8)	10.9 ± 1.2 (n=3)
15	1-(p-n-butylbenzyl) [MS-264]	491 (n=6)	14.6 ± 0.2 (n=2)
	1-(2,4-di-Me-benzyl)	404 (n=7)	10.2 ± 0.6 (n=3)
	1-(2,4,6-tri-Me-benzyl)	408 (n=7)	11.7 ± 0.6 (n=3)
	1,3-dibenzyl	254 (n=9)	8.40 (n=1)
	1,4-dibenzyl	378 (n=3)	8.66 (n=1)
20	1-(p-ethylbenzyl)-3-benzyl	nd	nd
	1,3-di(p-ethylbenzyl)	260 (n=3)	11.2 (n=1)
	1-(p-n-butylbenzyl)-3-benzyl	313 (n=3)	13.0 (n=1)

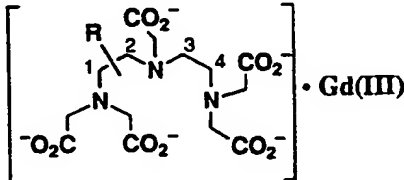
\* (Total number of animals in experiment listed as "n")

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Rabbit liver uptake

Anesthetized (pentobarbital, 35 mg/kg, ip.) adult rabbits (2.5-4 kg) were administered 50  $\mu\text{mol/kg}$  of a  $^{153}\text{Gd}$ -labeled chelate (approximately 30  $\mu\text{Ci/kg}$ ) via a cannulated marginal ear vein. Rabbits (1 per compound) were positioned supine under a gamma camera set with a photo peak energy of 100 keV  $\pm$  20% and equipped with a low energy parallel hole collimator. Immediately after chelate administration, dynamic planar whole body scintigraphy was performed every five minutes for at least one hour. Images were acquired in a 64 x 64 matrix, and region of interest (ROI) analysis was performed using the Image 1.52 software package (National Institutes of Health). Percent injected dose (% ID) in the liver at 5 min. intervals was calculated by dividing the counts in the liver ROI by the total counts in the body. Results are summarized in TABLE 3 and GRAPH 1.

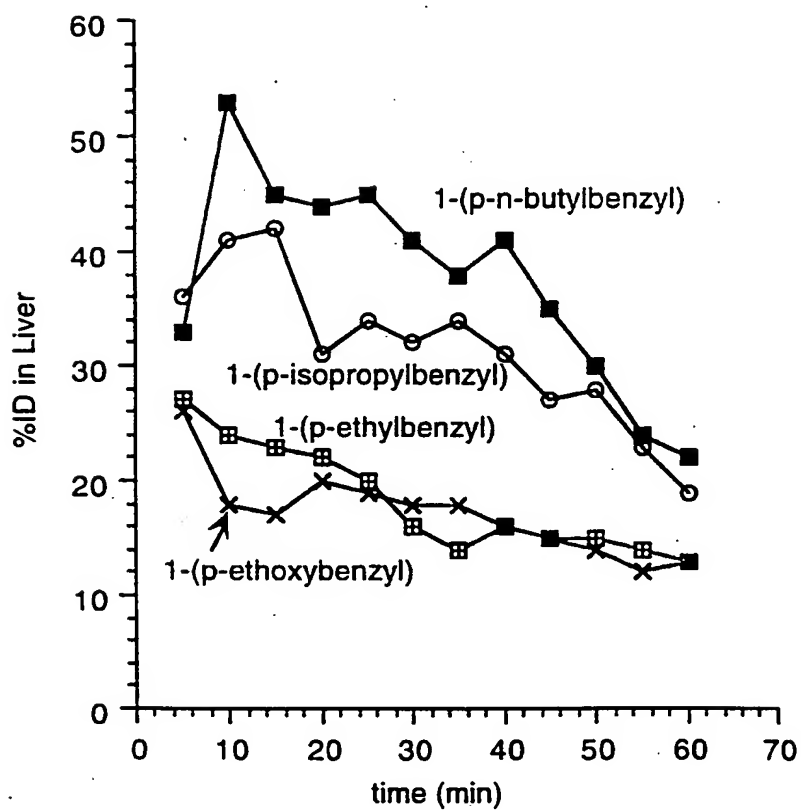
TABLE 3

20		Area under the % inj. dose in rabbit liver vs time curve (AUC) 0 to from 30 min (n=1 <u>rabbit per compd</u> )
25	<u>R</u>	
	1-(p-ethoxybenzyl) [EOB-DTPA]	480
	1-(p-ethylbenzyl)	553
	1-(p-isopropylbenzyl)	910
	1-(p-n-butylbenzyl) [MS-264]	1120

- 20 -

GRAPH 1

Rabbit Liver Scintigraphy of Gd153 Complexes  
(0.05mmol/kg)



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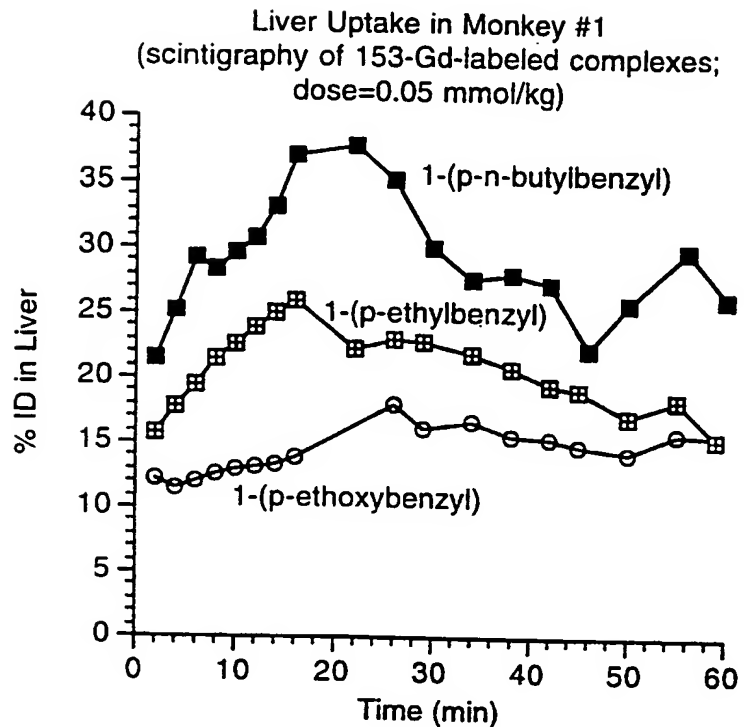
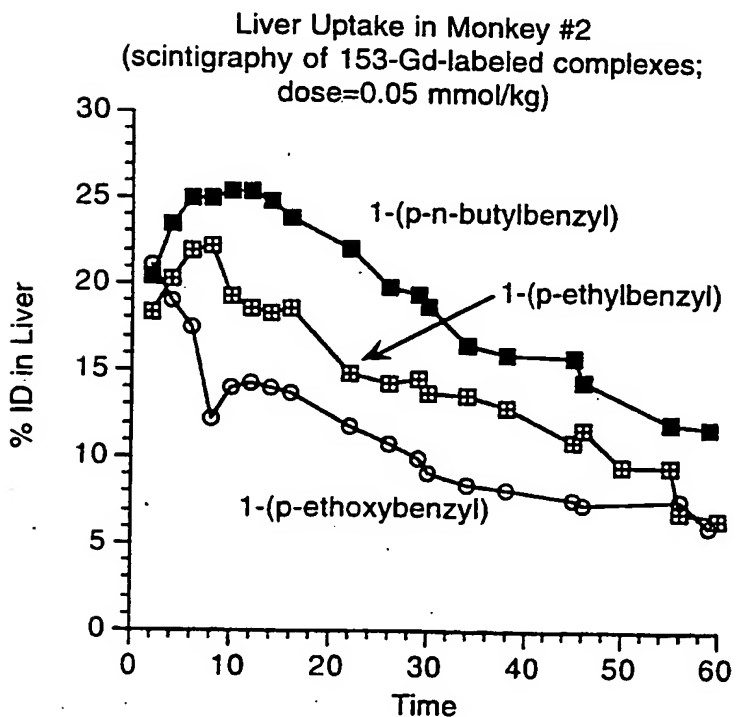
Monkey liver uptake

Two adult male cynomolgus monkeys were scintigraphically imaged after iv. administration of 50  $\mu\text{mol/kg}$  of the chelates labeled with  $^{153}\text{Gd}$  (approximately 40  $\mu\text{Ci/kg}$ ). All agents were given to each monkey on separate days. Monkeys were anesthetized with ketamine hydrochloride (i.m., to effect), intubated and maintained on isoflurane (0.5-2%) and positioned supine under a gamma camera prior to compound administration. Images were acquired and analyzed as described above. Results are summarized in TABLE 4 and GRAPHS 2 and 3.

TABLE 4

15		Area under the % inj. dose in monkey liver vs time curve (AUC) from 0 to 30 min	
	R	<u>Monkey # 1</u>	<u>Monkey #2</u>
20	1-(p-ethoxybenzyl) [EOB-DTPA]	396	380
	1-(p-ethylbenzyl)	617	494
	1-(p-n-butylbenzyl) [MS-264]	912	643

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GRAPH 2GRAPH 3





- 24 -

Synthesis of MS-264 $\alpha$ -Chloro- $\beta$ -p-n-butylphenylpropionitrile (2)

One and a half moles (237 mL) of 4-n-butylaniline was dissolved in acetone (1500 mL). The  
5 solution was cooled to 10 °C and water (150 mL) and concentrated aqueous HCl (300 mL) were added. Sodium nitrite (69.0 g, 1.73 moles) in water (200 mL) was added dropwise to the cooled solution. The solution was adjusted to pH 4 by addition of solid sodium  
10 acetate. Acrylonitrile (148 mL, 2.25 moles) was added, followed by  $\text{CuCl}_2 \cdot 2\text{H}_2\text{O}$  (38.4 g) in 50 mL  $\text{H}_2\text{O}$ . The reaction mixture was heated to 40°C and was stirred at this temperature for 3 h. The aqueous layer was extracted with ether. The combined organics were  
15 washed with water and saturated aqueous  $\text{NaHCO}_3$ , were dried over  $\text{Na}_2\text{SO}_4$ , filtered and concentrated under reduced pressure. The residue was fractionally distilled at reduced pressure. The product was collected at 115-130°C @ 2 mm Hg, as 166.3 g (50%  
20 yield) of an orange oil.

 $\alpha$ -Chloro- $\beta$ -p-n-butylphenylpropionic Acid (3)

The  $\alpha$ -chloro- $\beta$ -p-butylphenylpropionitrile 2 (166 g, 0.750 moles) was stirred over 240 mL formic acid and 160 mL concentrated HCl at reflux for 16 h.  
25 The reaction mixture was cooled, diluted with 300 mL water and washed with ethyl acetate. The ethyl acetate solution was washed with water, saturated aqueous NaCl, dried over  $\text{Na}_2\text{SO}_4$ , filtered and concentrated under reduced pressure. The bulk oil solidified on standing.  
30 The solid was recrystallized twice from hexane, yielding 139.0 g (0.58 moles, 77% yield) of a white solid, mp. 52.7-53.6°C.

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p-n-Butylphenylalanine (4)

$\alpha$ -Chloro- $\beta$ -p-butylphenylpropionic acid 3  
(128 g, 0.534 moles) was suspended in 800 mL 30%  
aqueous ammonia. The reaction was stirred for 4 days  
5 at 40°C in a tightly stoppered flask. After cooling to  
room temperature, the precipitate was removed by  
filtration through a coarse glass frit. The solid was  
washed thoroughly with ethanol followed by diethyl  
ether. The solid was dried under vacuum yielding  
10 54.3 g (0.245 moles, 46% yield) of p-butylphenylalanine  
as a white solid. The solid was recrystallized from 2N  
HCl to give a white solid, mp 218-223°C.

p-n-Butylphenylalanine Methyl Ester Hydrochloride (5)

Butylphenylalanine 4 (4.20 g, 19.0 mmoles)  
15 was suspended in 50 mL dry methanol. The solution was  
saturated with HCl gas and was refluxed for 2 hours.  
The solution was cooled and then added to 400 mL  
diethyl ether. The resulting solid was collected by  
vacuum filtration and was washed with ether. The solid  
20 was redissolved in a minimum volume of methanol and  
precipitated in ether as before resulting in 3.45 g  
(12.7 mmoles, 67% yield) of white solid (mp 164.5-  
165.1°C).

p-n-Butylphenylalanine Ethylenediamine Amide (6)

25 Butylphenylalanine methyl ester hydrochloride  
5 (3.45 g, 12.7 mmoles) was dissolved in 65 mL  
ethylenediamine and was stirred at room temperature for  
16 h. The ethylenediamine was removed by evaporation  
at reduced pressure. The residue was partitioned  
30 between 1 N NaOH and ethyl acetate and the aqueous was  
washed three times with ethyl acetate. The combined  
organic solutions were dried over Na<sub>2</sub>SO<sub>4</sub>, filtered and  
concentrated yielding 2.76 orange oil (83% yield). <sup>1</sup>H

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NMR (CDCl<sub>3</sub>)  $\delta$  7.50 (br. s, 1H), 7.15 (s, 4H), 3.60 (dd, 1H), 3.38-3.17 (3H), 2.85-2.50 (5H), 1.58 (m, 2H), 1.42-1.24 (6H), 0.93 (t, 3H).

5 1-p-n-Butylbenzyl-diethylenetriamine Trihydrochloride  
(7)

Butylphenylalanine ethylenediamine amid 6  
(2.76 g, 10.5 mmoles) was dissolved in 50 mL THF.  
Borane•THF (50 mL, 1.0 M) was added slowly to the  
stirred solution. The reaction was then refluxed under  
10 argon for 16 h, and quenched by careful addition of  
methanol. The reaction mixture was concentrated under  
reduced pressure. Dry ethanol (75 mL) was added and  
the solution was saturated with HCl gas at 0 °C. The  
mixture was brought to reflux for 24 h, after which the  
15 solution was cooled and concentrated. Ethyl ether was  
added to the concentrated solution and the resulting  
precipitate was collected by vacuum filtration and  
washed with ether, yielding 3.45 g (9.6 mmoles, 92%)  
white solid. Analysis calc'd for C<sub>15</sub>H<sub>27</sub>N<sub>3</sub>•3HCl: C,  
20 50.22; H, 8.43; N, 11.71; Cl, 29.64 found: C, 50.16;  
H, 8.44; N, 11.76; Cl, 29.57.

1-p-n-Butylbenzyl-diethylenetriamine penta-t-butyl  
Acetate (8)

Butylbenzyl-diethylenetriamine  
25 trihydrochloride 7 (3.45 g, 9.62 mmoles) was suspended  
in 50 mL dry DMF. Diisopropylethylamine (25 mL, 144  
mmoles) was added followed by 12.5 mL (77 mmoles) t-  
butylbromoacetate. The reaction was stirred at room  
temperature for 16 h under argon. The DMF and excess  
30 reagents were removed by evaporation under reduced  
pressure. The residue was partitioned between  
concentrated aqueous NaHCO<sub>3</sub> and chloroform. The  
organic solution was washed once each with water and  
saturated aqueous NaCl, dried over Na<sub>2</sub>SO<sub>4</sub>, filtered and

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concentrated under reduced pressure. The crude product was filtered through silica gel in 1:1 ethyl acetate:hexane. After being reconcentrated, the product was obtained as 6.88 g (8.39 mmol, 87% yield) of a light yellow oil.  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  7.15 (d, 2H), 7.04 (d, 2H), 3.50-3.35 (10H), 3.09 (m, 1H), 2.83 (m, 2H), 2.75-2.45 (8H), 1.65-1.10 (49H), 0.94 (t, 3H).

1-p-n-Butylbenzyl diethylenetriamine pentaacetic acid (9)

10 The penta-t-butyl ester **8** (6.88 g, 8.39 mmol) was dissolved in 1:1 dioxane:concentrated HCl and was stirred at room temperature for 16 h. The reaction mixture was concentrated under reduced pressure yielding the crude product as 4.79 g of a tan foam.

15 The product was eluted from a 10 g C-18 Sep-Pak cartridge with 40% aqueous acetonitrile. The acetonitrile was evaporated and the resulting aqueous solution was lyophilized producing 3.30 g (6.12 mmoles, 73% yield) of a flocculent white solid. Analysis

20 calc'd for  $\text{C}_{25}\text{H}_{37}\text{N}_3\text{O}_{10}$ : C, 55.65; H, 6.91; N, 7.79; found: C, 55.06; H, 6.99; N, 7.71; Cl, 0.00.

Gadolinium(III) 1-(p-n-butylbenzyl) diethylenetriamine pentaacetate dimeglumine salt (10)

Gadolinium oxide (1.99 g, 5.5 mmol), 1-(p-n-butylbenzyl) diethylenetriamine pentaacetic acid **9** (6.00 g, 10.0 mmol [90% by weight]) and N-methyl-D-glucamine (3.90 g, 20 mmol) were weighed in a 100-mL round-bottom flask and 30 mL of distilled water was added to it. The mixture was stirred at 95°C for six

25 hours. The resulting cloudy solution was loaded on a 10-g C-18 Sep-Pak® cartridge and the complex was eluted with water. The solvent was removed on a rotary evaporator, the resulting white solid residue was dried in high vacuo for 24 hours at 45°C. Yield: 9.78 g

30

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(88 %). Anal. Calcd for  $C_{39}H_{70}GdN_5O_{21}$ : C, 42.50; H, 6.40; N, 6.35; Gd, 14.27. Found: C, 42.11; H, 6.08; N, 6.35; Gd, 14.03.

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## CLAIMS:

1. A complex of a paramagnetic metal and 1-(p-n-butylbenzyl)diethylenetriamine pentaacetic acid and pharmaceutically acceptable salts thereof.
- 5           2. The complex according to claim 1, wherein the paramagnetic metal is selected from the group consisting of Gd(III), Fe(III), Mn(II), Mn(III), Cr(III), Cu(II), Dy(III), Tb(III), Ho(III), Er(III) and Eu(III).
- 10           3. The complex according to claim 1, wherein the paramagnetic metal is Gd(III).
4. Gadolinium(III)-1-(p-n-butylbenzyl)diethylenetriamine pentaacetate dimeglumine salt.
5. The complex according to claim 1 wherein  
15 1-(p-n-butylbenzyl)diethylenetriamine pentaacetic acid is present in racemic form.
6. The complex according to claim 1 wherein  
20 1-(p-n-butylbenzyl)diethylenetriamine pentaacetic acid is present enantiomerically enriched or enantiomerically pure form.
7. A pharmaceutical composition comprising a compound according to any one of claims 1 to 6 together with a pharmaceutically acceptable carrier, adjuvant or vehicle.
- 25           8. The pharmaceutical composition according to claim 7, further comprising a free organic ligand or a pharmaceutically acceptable salt thereof.

- 30 -

9. The pharmaceutical composition according to claim 7, further comprising a free organic ligand or calcium, sodium, meglumine or combination salts thereof.

5           10. The pharmaceutical composition according to claim 8 or claim 9, wherein the organic ligand is 1-(p-n-butylbenzyl)diethylenetriamine pentaacetic acid.

11. A method for tissue-specific contrast enhancement of MR images of mammalian organs and  
10 tissues comprising the step of administering a diagnostically effective amount of a complex according to any one of claims 1 to 6.

12. The method according to claim 11 for contrast enhancement of MR images of the hepatobiliary  
15 system.

13. Use of a compound according to any one of claims 1 to 6 for the manufacture of a MRI contrast agent.

14. A method for tissue-specific diagnostic  
20 X-ray imaging of mammalian organs and tissues comprising the step of administering a diagnostically effective amount of a complex according to any one of claims 1 to 6.



## INTERNATIONAL SEARCH REPORT

International Application No. PCT/US 95/04045

A. CLASSIFICATION OF SUBJECT MATTER  
A 61 K 49/00, A 61 K 49/04

According to International Patent Classification (IPC) or to both national classification and IPC <sup>6</sup>

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
A 61 K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP, A, 0 405 704 (SCHERING AKTIENGESELLSCHAFT) 02 January 1991 (02.01.91), abstract; claims. -----	1-14

☐ Further documents are listed in the continuation of box C.

☐ Patent family members are listed in annex.

## \* Special categories of cited documents:

- \*A\* document defining the general state of the art which is not considered to be of particular relevance
- \*E\* earlier document but published on or after the international filing date
- \*L\* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- \*O\* document referring to an oral disclosure, use, exhibition or other means
- \*P\* document published prior to the international filing date but later than the priority date claimed

\*T\* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

\*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

\*Y\* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

\*&\* document member of the same patent family

Date of the actual completion of the international search  
22 June 1995

Date of mailing of the international search report  
25. 07. 95

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Authorized officer

SCHNASS e.h.

# INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 95/04045

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 11, 12, 14  
because they relate to subject matter not required to be searched by this Authority, namely:  
  
Remark: Although claims 11, 12, 14 are directed to a method of treatment of the human and/or animal body by therapy (Rule 39.1(iv)PCT) the search was carried out and based on the alleged effects of the composition.
2. ☐ Claims Nos.:  
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

## ANHANG

zum internationalen Recherchen-  
bericht über die internationale  
Patentanmeldung Nr.

## ANNEX

to the International Search  
Report to the International Patent  
Application No.

## ANNEXE

au rapport de recherche inter-  
national relatif à la demande de brevet  
international n°

PCT/US 95/04045 SAE 108350

In diesem Anhang sind die Mitglieder  
der Patentfamilien der im obenge-  
nannten internationalen Recherchenbericht  
angeführten Patentdokumente angegeben.  
Diese Angaben dienen nur zur Unter-  
richtung und erfolgen ohne Gewähr.

This Annex lists the patent family  
members relating to the patent documents  
cited in the above-mentioned inter-  
national search report. The Office is  
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La présente annexe indique les  
membres de la famille de brevets  
relatifs aux documents de brevets cités  
dans le rapport de recherche inter-  
national visée ci-dessus. Les renseigne-  
ments fournis sont donnés à titre indica-  
tif et n'engagent pas la responsabilité  
de l'Office.

In Recherchenbericht angeführtes Patentdokument Patent document cited in search report Document de brevet cité dans le rapport de recherche	Datum der Veröffentlichung Publication date Date de publication	Mitglied(er) der Patentfamilie Patent family member(s) Membre(s) de la famille de brevets	Datum der Veröffentlichung Publication date Date de publication
EP A2 405704	02-01-91	AT E 115546	15-12-94
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		AU B2 637111	20-06-93
		CA AA 2020142	31-12-90
		CS A2 9003174	12-11-91
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		NO AO 902925	29-06-90
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		ZA A 9005129	24-04-91